"May be fatal or cause blindness if swallowed," and "Cannot be made nonpoisonous

- (3) If water is added at the time of manufacture, the label shall reflect the composition of the diluted product. If the addition of water reduces the methyl alcohol concentration to less than 4% by weight, the requirements of paragraph (c)(2) of this section shall not apply.
- (4) A back label shall be attached showing the word "ANTIDOTE", followed by suitable directions for an antidote.
- (d)(1) Reagent alcohol may be distributed in containers not exceeding 4 liters exclusively to laboratories or persons who require reagent alcohol for scientific use.
- (2) Reagent alcohol may be distributed in bulk containers to proprietors of bona fide laboratory supply houses for packaging and resale, and to any other person who was qualified to receive bulk shipments of reagent alcohol on the effective date of this part. Reagent alcohol may also be distributed in bulk containers to any person who has received approval of a letterhead application containing the following:
- (i) The applicant's name, address, and permit number, if any;
- (ii) A description of the security measures which will be taken to segregate reagent alcohol from denatured spirits or other alcohol which may be on the same premises:
- (iii) A statement that labels required by paragraph (c) of this section will be affixed to containers of reagent alcohol filled by the applicant;
- (iv) A statement that the applicant will allow appropriate TTB officers to inspect the applicant's premises; and
- (v) A statement that the applicant will comply with the requirements of §20.133.

(Approved by the Office of Management and Budget under control number 1512–0336)

[T.D. ATF- 199, 50 FR 1962, Mar. 6, 1985, as amended by T.D. ATF-435, 66 FR 5474, Jan. 19, 2001]

# § 20.118 Rubbing alcohol general-use formula.

(a) Rubbing alcohol is an article made with S.D.A. Formula No. 23–H (1)

containing 70% ethyl alcohol by volume (2) made in accordance with one of the two formulas prescribed in paragraph (b) of this section, and (3) labeled in accordance with §20.134(e) of this part.

(b) Either of the following two formulas is approved for manufacturing rubbing alcohol:

### Formula A

S.D.A. formula no. 23-H	103.3 fl. oz.
Sucrose octa-acetate	0.5 av.oz.
Water	q.s. 1 gal-
	lon

(If desired, ordorous, medicinal and/or colorative ingredients may be added.) Formula B

(If desired, odorous, medicinal and/or colorative ingredients may be added.)

### § 20.119 Toilet preparations containing not less than 10% essential oils general-use formula.

This general-use formula shall consist of an article containing not less than 10% essential oils by volume made with a formula of S.D.A. authorized for that article on the following list:

Article	Prod- uct code No.	Formula author- ized
Bath preparations	142	1, 3–A, 3–B, 3–C, 23–A, 30, 36, 38–B, 39–B, 39–C, 40, 40– A, 40–B, 40–C.
Colognes	122	38–B, 39, 39–A, 39–B, 39–C, 40, 40–A, 40– B, 40–C.
Deodorants (body)	114	23–A, 38–B, 39– B, 39–C, 40, 40–A, 40–B, 40–C.
Hair and scalp preparations	111	3–B, 23–A, 23–F, 23–H, 37, 38– B, 39, 39–A, 39–B, 39–C, 39–D, 40, 40– A, 40–B, 40–C.
Lotions and creams (body, face, and hand).	113	23–A, 23–H, 31– A, 37, 38–B, 39, 39–B, 39– C, 40, 40–A, 40–B, 40–C.
Perfume materials (processing)	121	38–B, 39, 39–B, 39–C, 40, 40– A, 40–B, 40–C.

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Article	Prod- uct code No.	Formula author- ized
Perfumes and perfume tinctures	121	38–B, 39, 39–B, 39–C, 40, 40– A, 40–B, 40–C.
Shampoos	141	1, 3–A, 3–B, 3–C, 23–A, 27–B, 31–A, 36, 38– B, 39–A, 39–B, 40, 40–A, 40– B, 40–C.
Soaps, toilet	142	1, 3–A, 3–C, 23– A, 30, 36, 38– B, 39–B, 39–C, 40, 40–A, 40– B, 40–C.
Toilet waters	122	38–B, 39, 39–A, 39–B, 39–C, 40, 40–A, 40– B, 40–C.

[T.D. ATF-199, 50 FR 9162, Mar. 6, 1985; 50 FR 20099, May 14, 1985]

# Subpart G—Requirements Relating to Articles

### § 20.131 Scope of subpart.

This subpart prescribes requirements relating to articles which may affect persons who are not required to obtain a permit under this part. These requirements, described in general terms §20.132, are imposed by law. Criminal penalties imposed for violating these requirements are described in §20.137. In this subpart, the term "article" means any substance or preparation in the manufacture of which denatured spirits are used, including the product obtained by further manufacture or by combination with other materials, if the article subjected to further manufacture or combination contained denatured spirits.

### § 20.132 General requirements.

- (a) Internal medicinal preparations and flavoring extracts—(1) Manufacture. No person shall use denatured spirits in the manufacture of medicinal preparations or flavoring extracts for internal human use where any of the spirits remain in the finished product.
- (2) Sale. No person shall sell or offer for sale for internal human use any medicinal preparations or flavoring extracts manufactured from denatured distilled spirits where any of the spirits remain in the finished product.

- (3) Labeling and advertising. Labeling and advertising of articles shall not imply that the article is intended for or suitable for internal human use.
- (b) Beverage use. No person shall sell or offer for sale any article containing denatured spirits for beverage purposes. Labeling and advertising of articles shall not imply that the article is intended for or suitable for use as a beverage.
- (c) Trafficking in articles. The appropriate TTB officer may impose the requirements of §20.133 on any person who reprocesses, rebottles, or repackages articles, deals in articles, or receives articles in containers exceeding one gallon.

### § 20.133 Registration of persons trafficking in articles.

- (a) Upon written notice from the appropriate TTB officer, any person who reprocesses, rebottles, or repackages articles, deals in articles, or receives articles in containers exceeding one gallon may be required to submit any of the following:
- (1) Nature of activities to be conducted;
- (2) Name and address of supplier;
- (3) Size and type of containers in which articles will be received and, if applicable, rebottled or repackaged;
- (4) Maximum quantity of each article to be obtained during any calendar month;
- (5) Description of the reprocessing operation;
- (6) Samples of the reprocessed article;
- (7) Labels and advertising materials; and,
- (8) Names and addresses of recipients of articles and quantities received;
- (b) The appropriate TTB officer shall prohibit any of the activities described in paragraph (a) of this section if the activities pose a jeopardy to the revenue, or a burden in administering this part.

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## § 20.134 Labeling.

(a) General. Except as provided in paragraph (b) or (c) of this section, each article shall, before removal from the manufacturer's premises, have a